

CP-1221

IN THE UNITED STATES PATENT & TRADEMARK OFFICE

PATENT APPLICATION

TITLE: PRIMARY INTERMEDIATE FOR OXIDATIVE COLORATION OF HAIR

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PRIMARY INTERMEDIATE FOR OXIDATIVE COLORATION OF HAIR

Field of the Invention

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This invention relates to a new compound, 4-amino-2-(1-hydroxy-ethyl)-phenol, and compositions containing this new compound as a primary intermediate for oxidative coloring of hair fibers.

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Background to the Invention

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Coloration of hair is a procedure practiced from antiquity employing a variety of means. In modern times, the method most extensively to color hair is an oxidative dyeing process utilizing one or more oxidative hair coloring agents in combination with one or more oxidizing agents.

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Most commonly a peroxy oxidizing agent is used in combination with one or more oxidative hair coloring agents, generally small molecules capable of diffusing into hair and comprising one or more primary intermediates and one or more couplers. In this procedure, a peroxide material, such as hydrogen peroxide, is employed to activate the small molecules of primary intermediates so that they react with couplers to form larger sized compounds in the hair shaft to color the hair in a variety of shades and colors.

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A wide variety of primary intermediates and couplers have been employed in such oxidative hair coloring systems and compositions. Among the primary intermediates employed there may be mentioned p-phenylenediamine, p-toluenediamine, p-aminophenol, 4-amino-3-methylphenol, N,N-bis(2-hydroxyethyl)-p-phenylenediamine, and 1-(2-hydroxyethyl)-4,5-diaminopyrazole and as couplers there may be mentioned

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resorcinol, 2-methylresorcinol, 3-aminophenol, 2,4-diaminophenoxyethanol, and 5-amino-2-methylphenol.

There are numerous additional requirements for oxidation dye compounds that are used to dye human hair besides the color or the desired intensity. Thus, the dye compounds must be unobjectionable in regard to toxicological and dermatological properties and must provide the desired hair color with a good light fastness, fastness to a permanent wave treatment, acid fastness and fastness to rubbing. The color of the hair dyed with the dye compounds in each case should be stable for at least 4 to 6 weeks to light, rubbing and chemical agents. Furthermore, an additional requirement is the production of a broad palette of different color shades using different developer and coupler substances. Many of the desired shades have been produced with dyes based on p-aminophenol. However, as indicated in U.S. Patent No. 4,997,451, the use of p-aminophenol is being questioned, for possible toxicological reasons. The proposed replacements for p-aminophenol have not proved entirely satisfactory. There is therefore a need for new primary intermediate compounds to meet one or more of the desired properties but not possessing the possible toxicological drawbacks possessed by p-aminophenol.

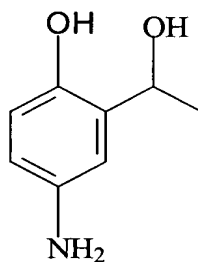
Summary of the Invention

It is therefore an object of this invention to provide new primary intermediate compound useful in place of p-aminophenol to provide a wide range of different color shades with various combinations of primary intermediates and couplers, but which avoids the drawback of p-aminophenol.

It has been discovered that the new compound 4-amino-2-(1-hydroxy-ethyl)-phenol is a suitable primary intermediate for hair coloring compositions and systems for providing good oxidative coloration of hair and for providing acceptable light fastness, fastness to shampooing, fastness to

permanent wave treatment, and suitable for providing a wide variety of different color shades with various primary intermediate and coupler compounds, but which avoids the drawback of p-aminophenol.

5 The invention provides the new compound of formula (1):



(1)

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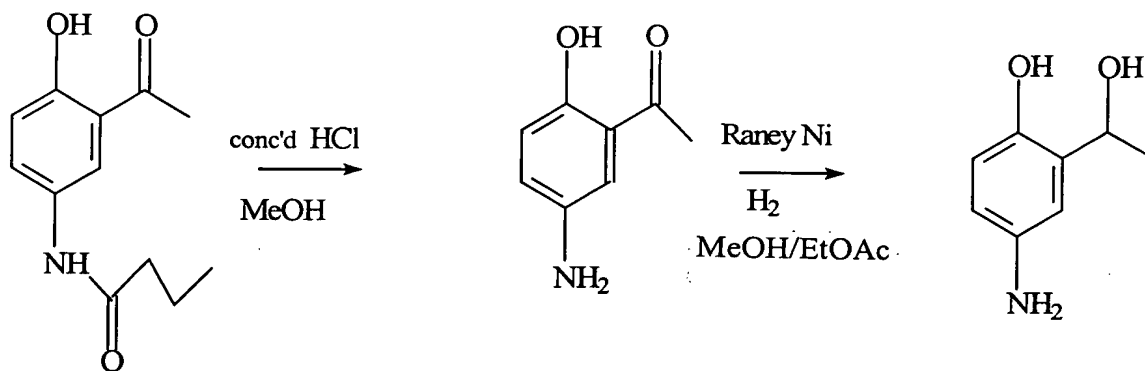
This novel primary intermediate is used to provide coloration to hair in which there is good dye uptake by the hair and provides shades or colors which are stable over a relatively long period of time. The novel primary intermediate provides for the dyeing of hair to impart color or shades that possess good wash fastness, have good selectivity, and do not undergo significant changes in coloration on exposure to light or perspiration.

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Detailed Description of the Invention

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The new 4-amino-2-(1-hydroxy-ethyl)-phenol compound of formula (1) of this invention can be prepared according to the following reaction sequence:



In this synthesis procedure 5-butylamido-2-hydroxyacetophenone is treated with a mixture of concentrated-HCl and methanol (MeOH) and neutralized with an alkali solution, e.g., NaOH solution, to precipitate 4-amino-2-hydroxyacetophenone, which upon hydrogenation with hydrogen and Raney Nickel in a MeOH/ethyl acetate (EtOAc) produces the 4-amino-2-(1-hydroxy-ethyl)-phenol compound of formula (1).

Synthesis Example 1

A suspension of 5-butylamido-2-hydroxyacetophenone (221.26 g, 1 mole) in a mixture of concentrated-HCl (500 mL) and MeOH (500 mL) was stirred at 80 °C for 24 h, cooled to room temperature and neutralized with 50% aqueous NaOH solution. The resulting precipitate was filtered, washed with cold water, and air-dried to produce 4-amino-2-hydroxyacetophenone (122.81 g, 81.2% yield): ¹HNMR (400 MHz, DMSO-d₆) δ 2.50 (s, 3H), 3.80 (bs, 3H), 6.69 (d, 1H, J=7.0 Hz), J=3 Hz), 6.86 (dd, 1H, J=7.0 Hz, J=3 Hz), 7.05 (1H,d, J=3 Hz); Ms *m/z* 181 (M⁺). Hydrogenation of 4-amino-2-hydroxyacetophenone (30.23 g, 200 mmole) in MeOH/EtOAc (150 mL/150 mL)) at 60 psi hydrogen pressure with wet Raney Nickel (20 g) for 24 h produced 4-amino-2-(1-hydroxy-ethyl)-phenol of formula (1) (24 g, 78.35 yield): ¹HNMR (400 MHz, DMSO-d₆) δ 1.23 (d, 3H), 4.35 (s, 2H), 4.88 (m, 2H), 6.26 (dd, 1H, J=8 Hz, J=3 Hz), 6.44 (d, 1H J=3 Hz), 6.60 (d, 1H, J=3 Hz), 8.27 (s, 1H); MS/*m/z* 153 (M⁺).

As used herein, the term "hair dyeing composition" (also synonymously referred to herein as the hair dye composition, the hair coloring composition, or the hair dye lotion) refers to the composition containing oxidation dyes, including the novel compound described herein, prior to admixture with the developer composition. The term "developer composition" (also referred to as the oxidizing agent composition or the peroxide composition) refers to compositions containing an oxidizing agent prior to admixture with the hair dyeing composition. The term "hair dye product" or "hair dye system" (also referred to as the hair dyeing system, hair dyeing product, or hair coloring system) interchangeably refer to the combination of the hair dyeing composition and the developer composition before admixture, and may further include a conditioner product and instructions, such product or system often being provided packaged as a kit. The term "hair dyeing product composition" refers to the composition formed by mixing the hair dyeing composition and the developer composition. "Carrier" (or vehicle or base) refers to the combination of ingredients contained in a composition excluding the active agents (e.g., the oxidation hair dyes of the hair dyeing composition).

Hair coloring (i.e., hair dyeing) compositions of this invention can contain, in combination with oxidation dye couplers, the novel primary intermediate of this invention as the sole primary intermediate or can also contain other primary intermediates. Thus, one or more suitable primary intermediates may be used in combination with the novel primary intermediate of this invention.

Suitable known primary intermediates include, for example,

p-phenylenediamine derivatives such as: benzene-1,4-diamine (commonly known as p-phenylenediamine), 2-methyl-benzene-1,4-diamine, 2-

chloro-benzene-1,4-diamine, N-phenyl-benzene-1,4-diamine, N-(2-ethoxyethyl)benzene-1,4-diamine, 2-[(4-amino-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol, (commonly known as N,N-bis(2-hydroxyethyl)-p-phenylenediamine) (2,5-diamino-phenyl)-methanol, 1-(2,5-diamino-phenyl)-ethanol, 2-(2,5-diamino-phenyl)-ethanol, N-(4-aminophenyl)benzene-1,4-diamine, 2,6-dimethyl-benzene-1,4-diamine, 2-isopropyl-benzene-1,4-diamine, 1-[(4-aminophenyl)amino]-propan-2-ol, 2-propyl-benzene-1,4-diamine, 1,3-bis[(4-aminophenyl)(2-hydroxyethyl)amino]propan-2-ol, N⁴,N⁴,2-trimethylbenzene-1,4-diamine, 2-methoxy-benzene-1,4-diamine, 1-(2,5-diaminophenyl)ethane-1,2-diol, 2,3-dimethyl-benzene-1,4-diamine, N-(4-amino-3-hydroxy-phenyl)-acetamide, 2,6-diethylbenzene-1,4-diamine, 2,5-dimethylbenzene-1,4-diamine, 2-thien-2-ylbenzene-1,4-diamine, 2-thien-3-ylbenzene-1,4-diamine, 2-pyridin-3-ylbenzene-1,4-diamine, 1,1'-biphenyl-2,5-diamine, 2-(methoxymethyl)benzene-1,4-diamine, 2-(aminomethyl)benzene-1,4-diamine, 2-(2,5-diaminophenoxy)ethanol, N-[2-(2,5-diaminophenoxy)ethyl]-acetamide, N,N-dimethylbenzene-1,4-diamine, N,N-diethylbenzene-1,4-diamine, N,N-dipropylbenzene-1,4-diamine, 2-[(4-aminophenyl)(ethyl)amino]ethanol, 2-[(4-amino-3-methyl-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol, N-(2-methoxyethyl)-benzene-1,4-diamine, 3-[(4-aminophenyl)amino]propan-1-ol, 3-[(4-aminophenyl)-amino]propane-1,2-diol, N-{4-[(4-aminophenyl)amino]butyl}benzene-1,4-diamine, and 2-[2-(2-{2-[(2,5-diaminophenyl)-oxy]ethoxy}ethoxy)ethoxy]benzene-1,4-diamine;

p-aminophenol derivatives such as: 4-amino-phenol (commonly known as p-aminophenol), 4-methylamino-phenol, 4-amino-3-methyl-phenol, 4-amino-2-hydroxymethyl-phenol, 4-amino-2-methyl-phenol, 4-amino-2-[(2-hydroxy-ethylamino)-methyl]-phenol, 4-amino-2-methoxymethyl-phenol, 5-amino-2-hydroxy-benzoic acid, 1-(5-amino-2-hydroxy-phenyl)-ethane-1,2-diol, 4-amino-2-(2-hydroxy-ethyl)-phenol, 4-amino-3-(hydroxymethyl)phenol, 4-amino-3-fluoro-phenol, 4-amino-2-(aminomethyl)-phenol, and 4-amino-2-fluoro-phenol;

o-aminophenol derivatives such as: 2-amino-phenol (commonly known as o-aminophenol), 2,4-diaminophenol, 2-amino-5-methyl-phenol, 2-amino-6-methyl-phenol, N-(4-amino-3-hydroxy-phenyl)-acetamide, and 2-amino-4-methyl-phenol; and

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heterocyclic derivatives such as: pyrimidine-2,4,5,6-tetramine (commonly known as 2,4,5,6-tetraaminopyridine), 1-methyl-1H-pyrazole-4,5-diamine, 2-(4,5-diamino-1H-pyrazol-1-yl)ethanol, N²,N²-dimethyl-pyridine-2,5-diamine, 2-[(3-amino-6-methoxypyridin-2-yl)amino]ethanol, 6-methoxy-N²-methyl-pyridine-2,3-diamine, 2,5,6-triaminopyrimidin-4(1H)-one, pyridine-2,5-diamine, 1-isopropyl-1H-pyrazole-4,5-diamine, 1-(4-methylbenzyl)-1H-pyrazole-4,5-diamine, 1-(benzyl)-1H-pyrazole-4,5-diamine, and 1-(4-chlorobenzyl)-1H-pyrazole-4,5-diamine.

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The novel primary intermediate of formula (1) of this invention may be used with any suitable coupler(s) in hair coloring compositions or systems of this invention.

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Suitable known couplers include, for example:

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phenols, resorcinol and naphthol derivatives such as: naphthalene-1,7-diol, benzene-1,3-diol, 4-chlorobenzene-1,3-diol, naphthalen-1-ol, 2-methyl-naphthalen-1-ol, naphthalene-1,5-diol, naphthalene-2,7-diol, benzene-1,4-diol, 2-methyl-benzene-1,3-diol, 7-amino-4-hydroxy-naphthalene-2-sulfonic acid, 2-isopropyl-5-methylphenol, 1,2,3,4-tetrahydro-naphthalene-1,5-diol, 2-chloro-benzene-1,3-diol, 4-hydroxy-naphthalene-1-sulfonic acid, benzene-1,2,3-triol, naphthalene-2,3-diol, 5-dichloro-2-methylbenzene-1,3-diol, 4,6-dichlorobenzene-1,3-diol, and 2,3-dihydroxy-[1,4]naphthoquinone;

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m-phenylenediamines such as: 2,4-diaminophenol, benzene-1,3-diamine, 2-(2,4-diamino-phenoxy)-ethanol, 2-[(3-amino-phenyl)-(2-

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hydroxy-ethyl)-amino]-ethanol, 2-methyl-benzene-1,3-diamine, 2-[[2-(2,4-diamino-phenoxy)-ethyl]-(2-hydroxy-ethyl)-amino]-ethanol, 4-{3-[(2,4-diaminophenyl)oxy]propoxy}benzene-1,3-diamine, 2-(2,4-diamino-phenyl)-ethanol, 2-(3-amino-4-methoxy-phenylamino)-ethanol, 4-(2-amino-ethoxy)-benzene-1,3-diamine, (2,4-diamino-phenoxy)-acetic acid, 2-[2,4-diamino-5-(2-hydroxy-ethoxy)-phenoxy]-ethanol, 4-ethoxy-6-methyl-benzene-1,3-diamine, 2-(2,4-diamino-5-methyl-phenoxy)-ethanol, 4,6-dimethoxy-benzene-1,3-diamine, 2-[3-(2-hydroxy-ethylamino)-2-methyl-phenylamino]-ethanol, 3-(2,4-diamino-phenoxy)-propan-1-ol, N-[3-(dimethylamino)phenyl]urea, 4-methoxy-6-methylbenzene-1,3-diamine, 4-fluoro-6-methylbenzene-1,3-diamine, 2-({3-[(2-hydroxyethyl)amino]-4,6-dimethoxyphenyl}-amino)ethanol, 3-(2,4-diaminophenoxy)-propane-1,2-diol, 2-[2-amino-4-(methylamino)-phenoxy]ethanol, 2-[(5-amino-2-ethoxy-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol, 2-[(3-aminophenyl)amino]ethanol, N-(2-aminoethyl)benzene-1,3-diamine, 4-[(2,4-diamino-phenyl)oxy]methoxy-benzene-1,3-diamine, and 2,4-dimethoxybenzene-1,3-diamine;

m-aminophenols such as: 3-amino-phenol, 2-(3-hydroxy-4-methyl-phenylamino)-acetamide, 2-(3-hydroxy-phenylamino)-acetamide, 5-amino-2-methyl-phenol, 5-(2-hydroxy-ethylamino)-2-methyl-phenol, 5-amino-2,4-dichloro-phenol, 3-amino-2-methyl-phenol, 3-amino-2-chloro-6-methyl-phenol, 5-amino-2-(2-hydroxy-ethoxy)-phenol, 2-chloro-5-(2,2,2-trifluoroethylamino)-phenol, 5-amino-4-chloro-2-methyl-phenol, 3-cyclopentylamino-phenol, 5-[(2-hydroxyethyl)amino]-4-methoxy-2-methylphenol, 5-amino-4-methoxy-2-methylphenol, 3-(dimethylamino)phenol, 3-(diethylamino)phenol, 5-amino-4-fluoro-2-methylphenol, 5-amino-4-ethoxy-2-methylphenol, 3-amino-2,4-dichloro-phenol, 3-[(2-methoxyethyl)amino]phenol, 3-[(2-hydroxyethyl)amino]phenol, 5-amino-2-ethyl-phenol, 5-amino-2-methoxyphenol, 5-[(3-hydroxypropyl)amino]-2-methylphenol, 3-[(3-hydroxy-2-methylphenyl)-amino]propane-1,2-diol, and 3-[(2-hydroxyethyl)amino]-2-methylphenol; and

heterocyclic derivatives such as: 3,4-dihydro-2H-1,4-benzoxazin-6-ol, 4-methyl-2-phenyl-2,4-dihydro-3H-pyrazol-3-one, 6-methoxyquinolin-8-amine, 4-methylpyridine-2,6-diol, 2,3-dihydro-1,4-benzodioxin-5-ol, 1,3-benzodioxol-5-ol, 2-(1,3-benzodioxol-5-ylamino)ethanol, 3,4-dimethylpyridine-2,6-diol, 5-chloropyridine-2,3-diol, 2,6-dimethoxypyridine-3,5-diamine, 1,3-benzodioxol-5-amine, 2-[[3,5-diamino-6-(2-hydroxy-ethoxy)-pyridin-2-yl]oxy]-ethanol, 1H-indol-4-ol, 5-amino-2,6-dimethoxypyridin-3-ol, 1H-indole-5,6-diol, 1H-indol-7-ol, 1H-indol-5-ol, 1H-indol-6-ol, 6-bromo-1,3-benzodioxol-5-ol, 2-aminopyridin-3-ol, pyridine-2,6-diamine, 3-[(3,5-diaminopyridin-2-yl)oxy]propane-1,2-diol, 5-[(3,5-diaminopyridin-2-yl)oxy]pentane-1,3-diol, 1H-indole-2,3-dione, indoline-5,6-diol, 3,5-dimethoxypyridine-2,6-diamine, 6-methoxypyridine-2,3-diamine, and 3,4-dihydro-2H-1,4-benzoxazin-6-amine.

Preferred primary intermediates include:

p-phenylenediamine derivatives such as: 2-methyl-benzene-1,4-diamine, benzene-1,4-diamine, 1-(2,5-diamino-phenyl)-ethanol, 2-(2,5-diamino-phenyl)-ethanol, N-(2-methoxyethyl)benzene-1,4-diamine, 2-[(4-amino-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol, and 1-(2,5-diaminophenyl)ethane-1,2-diol;

p-aminophenol derivatives such as 4-amino-phenol, 4-methylamino-phenol, 4-amino-3-methyl-phenol, 4-amino-2-methoxymethyl-phenol, and 1-(5-amino-2-hydroxy-phenyl)-ethane-1,2-diol;

o-aminophenol derivatives such as: 2-amino-phenol, 2-amino-5-methyl-phenol, 2-amino-6-methyl-phenol, N-(4-amino-3-hydroxy-phenyl)-acetamide, and 2-amino-4-methyl-phenol; and

heterocyclic derivatives such as: pyrimidine-2,4,5,6-tetramine, 1-methyl-1H-pyrazole-4,5-diamine, 2-(4,5-diamino-1H-pyrazol-1-yl)ethanol, 1-

(4-methylbenzyl)-1H-pyrazole-4,5-diamine, 1-(benzyl)-1H-pyrazole-4,5-diamine, and N²,N²-dimethyl-pyridine-2,5-diamine.

Preferred couplers include:

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phenols, resorcinol and naphthol derivatives such as: naphthalene-1,7-diol, benzene-1,3-diol, 4-chlorobenzene-1,3-diol, naphthalen-1-ol, 2-methyl-naphthalen-1-ol, naphthalene-1,5-diol, naphthalene-2,7-diol, benzene-1,4-diol, 2-methyl-benzene-1,3-diol, and 2-isopropyl-5-methylphenol;

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m-phenylenediamines such as: benzene-1,3-diamine, 2-(2,4-diamino-phenoxy)-ethanol, 4-{3-[(2,4-diaminophenyl)oxy]propoxy}benzene-1,3-diamine, 2-(3-amino-4-methoxy-phenylamino)-ethanol, 2-[2,4-diamino-5-(2-hydroxy-ethoxy)-phenoxy]-ethanol, and 3-(2,4-diamino-phenoxy)-propan-1-ol;

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m-aminophenols such as: 3-amino-phenol, 5-amino-2-methylphenol, 5-(2-hydroxy-ethylamino)-2-methylphenol, and 3-amino-2-methylphenol; and

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heterocyclic derivatives such as: 3,4-dihydro-2H-1,4-benzoxazin-6-ol, 4-methyl-2-phenyl-2,4-dihydro-3H-pyrazol-3-one, 1,3-benzodioxol-5-ol, 1,3-benzodioxol-5-amine, 1H-indol-4-ol, 1H-indole-5,6-diol, 1H-indol-7-ol, 1H-indol-5-ol, 1H-indol-6-ol, 1H-indole-2,3-dione, pyridine-2,6-diamine, and 2-aminopyridin-3-ol.

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Most preferred primary intermediates include:

p-phenylenediamine derivatives such as: 2-methyl-benzene-1,4-diamine, benzene-1,4-diamine, 2-(2,5-diamino-phenyl)-ethanol, 1-(2,5-diamino-phenyl)-ethanol, and 2-[(4-amino-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol;

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p-aminophenol derivatives such as: 4-amino-phenol, 4-methylamino-phenol, 4-amino-3-methyl-phenol, and 1-(5-amino-2-hydroxy-phenyl)-ethane-1,2-diol;

5 o-aminophenols such as: 2-amino-phenol, 2-amino-5-methyl-phenol, 2-amino-6-methyl-phenol, and N-(4-amino-3-hydroxy-phenyl)-acetamide; and

heterocyclic derivatives such as: pyrimidine-2,4,5,6-tetramine, 2-
10 (4,5-diamino-1H-pyrazol-1-yl)ethanol, 1-(4-methylbenzyl)-1H-pyrazole-4,5-diamine, and 1-(benzyl)-1H-pyrazole-4,5-diamine.

Most preferred couplers include:

15 phenols, resorcinol and naphthol derivatives such as: benzene-1,3-diol, 4-chlorobenzene-1,3-diol, naphthalen-1-ol, 2-methyl-naphthalen-1-ol, and 2-methyl-benzene-1,3-diol;

m-phenylenediamine such as: 2-(2,4-diamino-phenoxy)-ethanol,
20 2-(3-amino-4-methoxy-phenylamino)-ethanol, 2-[2,4-diamino-5-(2-hydroxy-ethoxy)-phenoxy]-ethanol, and 3-(2,4-diamino-phenoxy)-propan-1-ol;

m-aminophenols such as: 3-amino-phenol, 5-amino-2-methyl-phenol, 5-(2-hydroxy-ethylamino)-2-methyl-phenol, and 3-amino-2-methyl-
25 phenol; and

heterocyclic derivatives such as: 3,4-dihydro-2H-1,4-benzoxazin-6-ol, 4-methyl-2-phenyl-2,4-dihydro-3H-pyrazol-3-one, 1H-indol-6-ol, and 2-aminopyridin-3-ol.

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Understandably, the coupler compounds and the primary intermediate compounds, including the novel compound of the invention, in so far as they are bases, can be used as free bases or in the form of their

physiologically compatible salts with organic or inorganic acids, such as hydrochloric, citric, acetic, tartaric, or sulfuric acids, or, in so far as they have aromatic OH groups, in the form of their salts with bases, such as alkali phenolates.

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The total amount of dye precursors (e.g., primary intermediate and coupler compounds, including the novel compound of this invention) in the hair dyeing compositions of this invention is generally from about 0.002 to about 20, preferably from about 0.04 to about 10, and most preferably from about 0.1 to about 7.0 weight percent, based on the total weight of the hair dyeing composition. The primary intermediate and coupler compounds are generally used in molar equivalent amounts. However, it is possible to use the primary intermediate compounds in either excess or deficiency, i.e., a molar ratio of primary intermediate to coupler generally ranging from about 5:1 to about 1:5.

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The hair dyeing compositions of this invention will contain the primary intermediate of this invention in an effective dyeing amount, generally in an amount of from about 0.001 to about 10 weight percent by weight of the hair dye composition, preferably from about 0.01 to about 5.0 weight percent. Other primary intermediates, when present, are typically present in an amount such that in aggregate the concentration of primary intermediates in the composition is from about 0.002 to about 10 weight percent, preferably from about 0.01 to about 5.0 weight percent. The coupler(s) are present in an effective dyeing concentration, generally an amount of from about 0.001 to about 10.0 weight percent by weight of the hair dye composition, preferably from about 0.01 to about 5.0 weight percent. The remainder of the hair dye composition comprises a carrier or vehicle for the couplers and primary intermediates, and comprises various adjuvants as described below.

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Any suitable carrier or vehicle, generally an aqueous or hydroalcoholic solution, can be employed, preferably an aqueous solution.

The carrier or vehicle will generally comprise more than 80 weight percent of the hair dye composition, typically 90 to 99 weight percent, preferably 94 to 99 weight percent. The hair coloring compositions of this invention may contain as adjuvants one or more cationic, anionic, amphoteric, or zwitterionic surface active agents, perfumes, antioxidants such as ascorbic acid, thioglycolic acid or sodium sulfite, chelating and sequestering agents such as EDTA, thickening agents, alkalizing or acidifying agents, solvents, diluents, inerts, dispersing agents, penetrating agents, defoamers, enzymes, and other dye agents (e.g., synthetic direct and natural dyes). These adjuvants are cosmetic additive ingredients commonly used in compositions for coloring hair.

The hair dye compositions of the present invention are used by admixing them with a suitable oxidant, which reacts with the hair dye precursors to develop the hair dye. Any suitable oxidizing agent can be employed in the hair dye product compositions of this invention, particularly hydrogen peroxide (H_2O_2) or precursors therefor. Also suitable are urea peroxide, the alkali metal salts of persulfate, perborate, and percarbonate, especially the sodium salt, and melamine peroxide. The oxidant is usually provided in an aqueous composition generally referred to as the developer composition, which normally is provided as a separate component of the finished hair dye product and present in a separate container. The developer composition may also contain, to the extent compatible, various ingredients needed to form the developer composition, i.e., peroxide stabilizers, foam formers, etc., and may incorporate one or more of the adjuvants referred to above, e.g., surface active agents, thickeners, pH modifiers, etc. Upon mixing the hair coloring composition and the developer composition to form a hair dye product composition, the adjuvants are provided in the hair dye product composition as it is applied to the hair to achieve desired product attributes, e.g., pH, viscosity, rheology, etc.

The form of the hair dye product compositions according to the invention can be, for example, a solution, especially an aqueous or aqueous-

alcoholic solution. However, the form that is preferred is a thick liquid, cream, gel or an emulsion whose composition is a mixture of the dye ingredients with the conventional cosmetic additive ingredients suitable for the particular preparation.

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Suitable conventional cosmetic additive ingredients useful in the hair dye and developer compositions, and hence in the hair dye product compositions of this invention are described below, and may be used to obtain desired characteristics of the hair dye, developer and hair dye product compositions.

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Solvents: In addition to water, solvents that can be used are lower alkanols (e.g., ethanol, propanol, isopropanol, benzyl alcohol); polyols (e.g., carbitols, propylene glycol, hexylene glycol, glycerin). See WO 98/27941 (section on diluents) incorporated by reference. See also US 6027538 incorporated by reference. Under suitable processing, higher alcohols, such as C8 to C18 fatty alcohols, especially cetyl alcohol, are suitable organic solvents, provided they are first liquified by melting, typically at low temperature (50 to 80 °C), before incorporation of other, usually lipophilic, materials.

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The organic solvents are typically present in the hair dye compositions in an amount of from about 5 to about 30% by weight of the hair dye composition. Water is usually present in an amount of from about 5 to about 90% by weight of the hair dye composition, preferably from about 15 to about 75% by weight and most preferably from about 30 to about 65% by weight.

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Surfactants: These materials are from the classes of anionic, cationic, amphoteric (including zwitterionic surfactants) or nonionic surfactant compounds. (Cationic surfactants, generally included as hair conditioning materials, are considered separately below.) Suitable surfactants, other than cationic surfactants, include fatty alcohol sulfates, ethoxylated fatty alcohol

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sulfates, alkylsulfonates, alkylbenzensulfonates, alkyltrimethylammonium salts, alkylbetaines, ethoxylated fatty alcohols, ethoxylated fatty acids, ethoxylated alkylphenols, block polymers of ethylene and/orpropylene glycol, glycerol esters, phosphate esters, fatty acid alkanol amides and ethoxylated fatty acid esters, alkyl sulfates, ethoxylated alkyl sulfates, alkyl glyceryl ether sulfonates, methyl acyl taurates, acyl isethionates, alkyl ethoxy carboxylates, fatty acid mono- and diethanolamides. Especially useful are sodium and ammonium alkyl sulfates, sodium and ammonium ether sulfates having 1 to 3 ethylene oxide groups, and nonionic surfactants sold as Tergitols, e.g., C11-C15 Pareth-9, and Neodols, e.g., C12-C15 Pareth-3. They are included for various reasons, e.g., to assist in thickening, for forming emulsions, to help in wetting hair during application of the hair dye product composition, etc. Amphoteric surfactants include, for example, the asparagine derivatives as well betaines, sultaines, glycinate and propionates having an alkyl or alkylamido group of from about 10 to about 20 carbon atoms. Typical amphoteric surfactants suitable for use in this invention include lauryl betaine, lauroamphoglycinate, lauroamphopropionate, lauryl sultaine, myristamidopropyl betaine, myristyl betaine, stearoamphopropylsulfonate, cocamidoethyl betaine, cocamidopropyl betaine, cocoamphoglycinate, cocoamphocarboxypropionate, cocoamphocarboxyglycinate, cocobetaine, and cocoamphopropionate. Reference is made to WO 98/52523 published November 26, 1998 and WO 01/62221 published August 30, 2001, both incorporated herein by reference thereto.

The amount of surfactants in the hair dye compositions is normally from about 0.1% to 30% by weight, preferably 1% to 15% by weight.

Thickeners: Suitable thickeners include such as higher fatty alcohols, starches, cellulose derivatives, petrolatum, paraffin oil, fatty acids and anionic and nonionic polymeric thickeners based on polyacrylic and polyurethane polymers. Examples are hydroxyethyl cellulose, hydroxymethylcellulose and other cellulose derivatives, hydrophobically modified anionic polymers and

nonionic polymers, particularly such polymers having both hydrophilic and hydrophobic moieties (i.e., amphiphilic polymers). Useful nonionic polymers include polyurethane derivatives such as PEG-150/stearyl alcohol/SDMI copolymer. Suitable polyether urethanes are Aculyn® 22, 44 and Aculyn® 46
 5 polymers sold by Rohm & Haas. Other useful amphiphilic polymers are disclosed in US Pat. No. 6010541 incorporated by reference. See also WO 01/62221 mentioned above. Examples of anionic polymers that can be used as thickeners are acrylates copolymer, acrylates/ceteth-20 methacrylates copolymer, acrylates/ceteth-20 itaconate copolymer, and acrylates/beheneth-
 10 25 acrylates copolymers. In the case of the associative type of thickeners, e.g., Aculyns 22, 44 and 46, the polymer may be included in one of either the hair dye composition or the developer composition of the hair dye product and the surfactant material in the another. Thus, upon mixing of the hair dye and developer compositions, the requisite viscosity is obtained. The thickeners are
 15 provided in an amount to provide a suitably thick product as it is applied to the hair. Such products generally have a viscosity of from 1000 to 100000 cps, and often have a thixotropic rheology.

pH Modifying agents: Suitable materials that are used to adjust pH of the hair
 20 dye compositions include alkalizers such alkali metal and ammonium hydroxides and carbonates, especially sodium hydroxide and ammonium carbonate, ammonia, organic amines including methylethanolamine, aminomethylpropanol, mono-, di-, and triethanolamine, and acidulents such as inorganic and inorganic acids, for example phosphoric acid, acetic acid,
 25 ascorbic acid, citric acid or tartaric acid, hydrochloric acid, etc. See US patent 6027538 incorporated by reference.

Conditioners: Suitable materials include silicones and silicone derivatives; hydrocarbon oils; monomeric quaternary compounds, and quaternized
 30 polymers. Monomeric quaternary compounds are typically cationic compounds, but may also include betaines and other amphoteric and zwitterionic materials that provide a conditioning effect. Suitable monomeric

quaternary compounds include behentrialkonium chloride, behentrimonium chloride, benzalkonium bromide or chloride, benzyl triethyl ammonium chloride, bis-hydroxyethyl tallowmonium chloride, C12-18 dialkyldimonium chloride, cetalkonium chloride, cetearrimonium bromide and chloride, 5 cetrimonium bromide, chloride and methosulfate, cetylpyridonium chloride, cocamidopropyl ethyldimonium ethosulfate, cocamidopropyl ethosulfate, coco-ethyldimonium ethosulfate, cocotrimonium chloride and ethosulfate, dibehenyl dimonium chloride, dicetyldimonium chloride, dicocodimonium chloride, dilauryl dimonium chloride, disoydimonium chloride, 10 ditallowdimonium chloride, hydrogenated tallow trimonium chloride, hydroxyethyl cetyl dimonium chloride, myristalkonium chloride, olealkonium chloride, soyethomonium ethosulfate, soytrimonium chloride, stearalkonium chloride, and many other compounds. See WO 98/27941 incorporated by reference. Quaternized polymers are typically cationic polymers, but may 15 also include amphoteric and zwitterionic polymers. Useful polymers are exemplified by polyquaternium-4, polyquaternium-6, polyquaternium-7, polyquaternium-8, polyquaternium-9, polyquaternium-10, polyquaternium-22, polyquaternium-32, polyquaternium-39, polyquaternium-44 and polyquaternium-47. Silicones suitable to condition hair are dimethicone, 20 amodimethicone, dimethicone copolyol and dimethiconol. See also WO 99/34770 published July 15, 1999, incorporated by reference, for suitable silicones. Suitable hydrocarbon oils would include mineral oil.

Conditioners are usually present in the hair dye composition in an 25 amount of from about 0.01 to about 5% by weight of the composition.

Direct Dyes: The hair dyeing compositions according to the invention can also contain compatible direct dyes including Disperse Black 9, HC Yellow 2, HC Yellow 4, HC Yellow 15, 4-nitro-o-phenylenediamine, 2-amino-6-chloro-4- 30 nitrophenol, HC Red 3, Disperse Violet 1, HC Blue 2, Disperse Blue 3, and Disperse Blue 377. These direct dyes can be contained in the hair coloring

compositions of the invention in an amount of from about 0.05 to 4.0 percent by weight.

5 Natural ingredients: For example, proteins and protein derivatives, and plant materials such as aloe, chamomile and henna extracts.

Other adjuvants include polysaccharides, alkylpolyglycosides, buffers, chelating and sequestrant agents, antioxidants, and peroxide stabilizing agents as mentioned in WO 01/62221, etc.

10

The adjuvants referred to above but not specifically identified that are suitable are listed in the International Cosmetics Ingredient Dictionary and Handbook, (Eighth Edition) published by The Cosmetics, Toiletry, and Fragrance Association, incorporated by reference. In particular reference is made to Volume 2, Section 3 (Chemical Classes) and Section 4 (Functions) are useful in identifying a specific adjuvant to achieve a particular purpose or multipurpose.

15

The above-mentioned conventional cosmetic ingredients are used in amounts suitable for their functional purposes. For example, the surfactants used as wetting agents, associative agents, and emulsifiers are generally present in concentrations of from about 0.1 to 30 percent by weight, the thickeners are useful in an amount of from about 0.1 to 25 percent by weight, and the hair care functional materials are typically used in concentrations of from about 0.01 to 5.0 percent by weight.

20

25

The hair dyeing product composition as it is applied to the hair, i.e., after mixing the hair dye composition according to the invention and the developer, can be weakly acidic, neutral or alkaline according to their composition. The hair dye compositions can have pH values of from about 6 to 11.5, preferably from about 6.8 to about 10, and especially from about 8 to about 10. The pH of the developer composition is typically acidic, and generally the pH is from about 2.5 to about 6.5, usually about 3 to 5. The pH

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of the hair dye and developer compositions is adjusted using a pH modifier as mentioned above.

In order to use the hair coloring composition for dyeing hair, the above-described hair coloring compositions according to the invention are mixed with an oxidizing agent immediately prior to use and a sufficient amount of the mixture is applied to the hair, according to the hair abundance, generally from about 60 to 200 grams. Some of the adjuvants listed above (e.g., thickeners, conditioners, etc.) can be provided in the dye composition or the developer, or both, depending on the nature of the ingredients, possible interactions, etc., as is well known in the art.

Typically, hydrogen peroxide, or its addition compounds with urea, melamine, sodium borate or sodium carbonate, can be used in the form of a 3 to 12 percent, preferably 6 percent, aqueous solution as the oxidizing agent for developing the hair dye. Oxygen can also be used as the oxidizing agent. If a 6 percent hydrogen peroxide solution is used as oxidizing agent, the weight ratio of hair coloring composition and developer composition is 5:1 to 1:5, but preferably 1:1. In general, the hair dyeing composition comprising primary intermediate(s) and coupler(s), including at least one of the compounds of formula (1), is prepared and then, at the time of use, the oxidizing agents, such as H_2O_2 , contained in a developer composition is admixed therewith until an essentially homogenous composition is obtained, which is applied shortly after preparation to the hair to be dyed and permitted to remain in contact with the hair for a dyeing effective amount of time. The mixture of the oxidizing agent and the dye composition of the invention (i.e., the hair dye product composition) is allowed to act on the hair for about 2 to about 60 minutes, preferably about 15 to 45, especially about 30 minutes, at about 15 to 50°C, the hair is rinsed with water, and dried. If necessary, it is washed with a shampoo and rinsed, e.g., with water or a weakly acidic solution, such as a citric acid or tartaric acid solution. Subsequently the hair is dried. Optionally, a separate conditioning product may also be provided.

Together the hair dye composition of the present invention comprising the hair dye primary intermediate (1) and the developer composition comprising the oxidizing agent form a system for dyeing hair.

5 This system may be provided as a kit comprising in a single package separate containers of the hair dye composition, the developer, the optional conditioner or other hair treatment product, and instructions for use.

Especially useful primary intermediates of formula (1) of this

10 invention will provide hair coloring compositions having outstanding color fastness, especially light fastness, fastness to washing and fastness to rubbing.

Dyeing Example 1

15 The following composition shown in Table 1 can be used for dyeing Piedmont hair. 100 g of the dyeing composition is mixed with 100 g 20 volume hydrogen peroxide. The resulting mixture is applied to the hair and permitted to remain in contact with the hair for 30 minutes. The dyed hair

20 is then shampooed, rinsed with water and dried. The ranges of ingredients set out in Table 1 are illustrative of useful concentrations of the recited materials in a hair dye product.

TABLE 1
Composition for Dyeing Hair

Ingredients	Range (wt %)	Weight (%)
Cocamidopropyl betaine	0-25	17.00
Polyquaternium-22	0-7	5.00
Monoethanolamine ¹	0-15	2.00
Oleic Acid	2-22	0.75
Citric Acid	0-3	0.10
28% Ammonium hydroxide ¹	0-15	5.00
Behentrimonium chloride	1-5	0.50
Sodium sulfite	0-1	0.10
EDTA	0-1	0.10
Erythorbic acid	0-1	0.40
Ethoxydiglycol	1-10	3.50
C11-15 Pareth-9 (Tergitol 15-S-9)	0.5-5	1.00
C12-15 Pareth-3 (Neodol 25-3)	0.25-5	0.50
Isopropanol	2-10	4.00
Propylene glycol	1-12	2.00
p-phenylenediamine	0-5	1 mmole
N,N-Bis(hydroxyethyl)-p-phenylene diamine	0-5	1 mmole
3-Methyl-p-aminophenol	0-5	1 mmole
p-Aminophenol	0-5	1 mmole
Primary Intermediate of this invention	0.5-5	4 mmoles
5-Amino-2-Methyl Phenol ²	0-5	3 mmoles
2,4-Diaminophenoxyethanol ²	0-5	3 mmoles
M-Phenylenediamine ²	0-5	1 mmole
Water	qs to 100.00	qs to 100.00

¹ In the aggregate, these ingredients are in the range of 2 to 15% by weight.

² At least one of these dye precursors is typically present.

Dyeing Example 2

The hair dye composition in the table below was used for dyeing Piedmont hair weighing from 700 to 900 mg. 1 mL of the hair dye composition was mixed with 1 mL developer solution comprising 50% hydrogen peroxide 12.0 g, ACULYN® 22 1.0 g, ACULYN® 33 12.0 g, phosphoric acid to pH 3.0 and water qs to 100 g. The resulting mixture was applied to the hair tresses mounted on a glass plate and then stored at 40 °C

for 30 minutes, washed, shampooed, and dried. Color was evaluated using a Minolta Spectrophotometer CM-3700d. Color space is CIE L*a*b* and illuminant is D65 daylight with 10° observer. The color space, L* indicates lightness and a* and b* are the chromaticity coordinates. +a* is the red direction, -a* is the green direction, +b* is the yellow direction and -b* is the blue direction.

Composition for the Hair Dye Composition

Ingredients	Weight (%)
Cocamidopropyl betaine	17.00
Monoethanolamine	2.00
Oleic Acid	0.75
Citric Acid	0.10
28%Ammonia	5.00
Behentrimonium chloride	0.50
Sodium sulfite	0.10
EDTA	0.10
Erythorbic acid	0.40
Ethoxydiglycol	3.50
C11-15 Pareth-9 (Tergitol 15-S-9)	1.00
C12-15 Pareth-3 (Neodol 25-3)	0.50
Isopropanol	4.00
Propylene glycol	2.00
Coupler of Table 2	0.025 M
Primary intermediate of formula (1)	0.025 M
Water	qs to 100.00

10

Table 2 provides the results of the dyeing tests using the novel primary intermediate 1 of this invention, and also includes results for comparative dye compositions.

Table 2. Comparative color results with PAP derivatives				
Primary Int.	Coupler	L*	a*	b*
1	2,4-Diaminophenoxyethanol	47.9	10.1	7.5
3-Me-PAP	2,4-Diaminophenoxyethanol	51.2	7.9	5.4
2-Me-PAP	2,4-Diaminophenoxyethanol	55.9	5.2	10.9
1	5-Amino-2-methylphenol	63.6	13.6	22.7
3-MePAP	5-Amino-2-methylphenol	65.1	14.4	21.6
2-Me-PAP	5-Amino-2-methylphenol	68.6	8	21.7
1	1-Naphthol	65.5	12	16.5
3-MePAP	1-Naphthol	67	8.9	16.2
2-Me-PAP	1-Naphthol	67.8	8	17.7
1	2-Methyl-1-naphthol	63.7	16.5	16.3
3-MePAP	2-Methyl-1-naphthol	63	15	12.5
2-Me-PAP	2-Methyl-1-naphthol	64	15.3	14.6

- 5 1= 4-amino-2-(1-hydroxy-ethyl)-phenol of formula (1) of this invention
 3-Me-PAP= 3-methyl-p-aminophenol
 2-Me-PAP= 2-methyl-p-aminophenol

10 Surprisingly, the color obtained from coupling 4-amino-2-(1-hydroxy-ethyl)-phenol with 5-amino-2-methylphenol is redder than from coupling 2-methyl-p-aminophenol with 5-amino-2-methylphenol. The +a* value, which indicates redness, has a 13.6 value for 4-amino-2-(1-hydroxy-ethyl)-phenol compared to 8.0 for 2-methyl-p-aminophenol. Also, coupling of 4-amino-2-(1-hydroxy-ethyl)-phenol with 1-naphthol produces redder color than 2-methyl-p-aminophenol with 1-naphthol. The a* value for 4-amino-2-(1-hydroxy-ethyl)-phenol was 12 which was higher than the value of 8 for 2-methyl-p-aminophenol. Coloring intensity obtained from coupling 4-amino-2-(1-hydroxy-ethyl)-phenol with various couplers is in general higher than that obtained from 2-methyl-p-aminophenol and 3-methyl-p-aminophenol.

20

Dyeing Examples 3 to 6

25 100g of the following hair dye compositions shown in Tables 3 and 4 were mixed with 100 g of 20 volume hydrogen peroxide. The resulting mixture was applied on Piedmont hair and permitted to remain in contact with

at room temperature. Thus dyed hair was then shampooed and rinsed with water and dried.

5

**Table 3. Dyeing Compositions of Orange Red and Dark Red
(Examples 3 and 4)**

	Composition A (%)	Composition B (%)
Cocamidopropyl betaine	17.00	17.00
Ethanolamine	2	2
Oleic Acid	0.75	0.75
Citric Acid	0.1	0.1
Ammonium hydroxide	5.0	5.0
Behentrimonium chloride	0.5	0.5
Sodium sulfite	0.1	0.1
EDTA	0.1	0.1
Erythorbic acid	0.4	0.4
4-amino-2-(1-hydroxy-ethyl)-phenol	0.153	0.153
N-(2-Hydroxyethyl)-4,5-diaminopyrazole sulfate	0.240	
p-Phenylenediamine		0.108
2-Methyl-5-aminophenol	0.246	0.246
Water	QS 100	QS 100
Shade on gray hair	Orange Red	Dark Red

For the comparative purpose, the compound 4-amino-2-(1-hydroxy-ethyl)-phenol of Table 3 compositions A and B was replaced with equal molar 3-methyl-p-aminophenol and are Comparative Compositions C and D in Table 4. The coloration results are set forth in Table 4.

10

**Table 4. Compositions containing 3-methyl-p-aminophenol
(Comparative Examples 5 and 6)**

	Composition C (%)	Composition D (%)
Cocamidopropyl betaine	17.00	17.00
Ethanolamine	2	2
Oleic Acid	0.75	0.75
Citric Acid	0.1	0.1
Ammonium hydroxide	5.0	5.0
Behentrimonium chloride	0.5	0.5
Sodium sulfite	0.1	0.1
EDTA	0.1	0.1
Erythorbic acid	0.4	0.4
3-Methyl-p-aminophenol	0.123	0.123
N-(2-Hydroxyethyl)-4,5-diaminopyrazole sulfate	0.240	
p-Phenylenediamine		0.108
2-Methyl-5-aminophenol	0.246	0.246
Water	QS 100	QS 100
Shade on gray hair	Orange Red	Dark Red

5 The CIE L*a*b* values from the composition A which contains the compound 4-amino-2-(1-hydroxy-ethyl)-phenol are very similar to those from the composition C which contains 3-methyl-p-aminophenol. The composition B and D also give similar CIE L*a*b* values (Table 5).

Table 5. The CIE L*a*b* values obtained from Composition A, B, C and D

Composition	L*	a*	b*
A	52.49	35.85	30.63
B	39.08	14.84	4.86
C	52.23	37.99	31.65
D	37.41	15.86	4.78

15 The compound 4-amino-2-(1-hydroxy-ethyl)-phenol imparts very similar coloration to that of 3-methyl-p-aminophenol in the presence of other primary intermediates and couplers as shown in the results in Table 5. Both Composition A containing 4-amino-2-(1-hydroxy-ethyl)-phenol and Composition C having 3-methyl-p-aminophenol together with a pyrazole derivative and 5-amino-2-methylphenol color Piedmont hair almost identical orange red. Similar results are obtained with Compositions B and D. Overall
20 colors generated with 4-amino-2-(1-hydroxy-ethyl)-phenol are similar to those

with 3-methyl-p-aminophenol as shown by Tables 2 and 5. These results are surprising since 4-amino-2-(1-hydroxy-ethyl)-phenol is a 2-substituted p-aminophenol derivative as opposed to 3-methyl-p-aminophenol which is a 3-substituted p-aminophenol derivative.

5

Exemplary combinations of hair coloring components employing the compound of formula (1) of this invention are shown in combinations C1 to C136 in Tables A through H. Reading down the columns in Table A, the Xes demonstrate combinations of dyes that can be formulated according to the present invention. For example, in Combination No. C1 in Column 4 of Table A, the 4-amino-2-(1-hydroxy-ethyl)-phenol primary intermediate compound of formula (1) of this invention (Row 1 of Table A), can be combined with with 2-amino-phenol.

10
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Table A. Dye Combinations

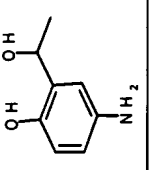
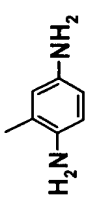
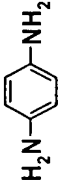

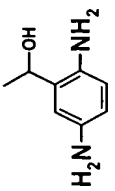
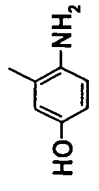
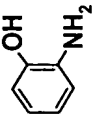
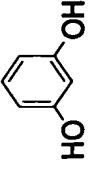
Structure	IUPAC Name	Name	C1	C2	C3	C4	C5	C6	C7	C8	C9	C10	C11
	4-Amino-2-(1-hydroxyethyl)-phenol		X	X	X	X	X	X	X	X	X	X	X
	2-Methylbenzene-1,4-diamine	p-Toluene-diamine											
	Benzene-1,4-diamine	p-Phenylene-diamine											
	2-[(4-Amino-phenyl)-(2-hydroxy-ethyl)-amino]-ethanol	N,N-Bis(2-hydroxyethyl)-p-phenylene-diamine											
	1-(2,5-Diamino-phenyl)-ethanol	1-Hydroxyethyl-p-phenylenediamine											
	4-Amino-3-methylphenol	3-Methyl-p-aminophenol											
	2-Amino-phenol	o-Aminophenol	X									X	
	Benzene-1,3-diol	Resorcinol		X								X	X

Table A (continued). Dye Combinations

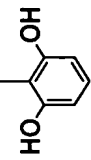
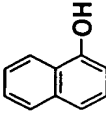
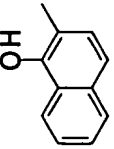
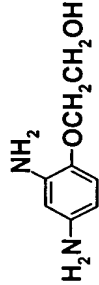
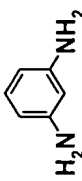
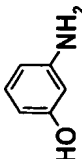
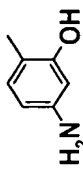
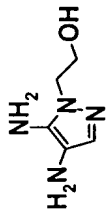
Structure	IUPAC Name	Name	C1	C2	C3	C4	C5	C6	C7	C8	C9	C10	C11
	2-Methylbenzene-1,3-diol	2-Methyl-resorcinol			x								x
	Naphthalen-1-ol	1-Naphthol				x							
	2-Methyl-naphthalen-1-ol	2-Methyl-1-naphthol					x						
	2-(2,4-Diaminophenoxy)ethanol	2,4-Diaminophenoxyethanol						x					
	Benzene-1,3-diamine	m-Phenylenediamine							x				
	3-Amino-phenol	m-Aminophenol								x			
	5-Amino-2-methylphenol	2-Hydroxy-4-aminotoluene									x		
	2-(4,5-Diaminopyrazol-1-yl)ethanol	1-Hydroxyethyl-4,5-diamino-pyrazole											

Table B. Dye Combinations

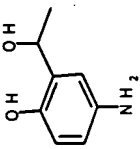
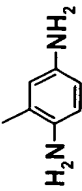
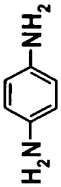

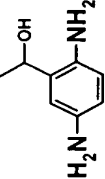
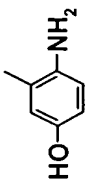
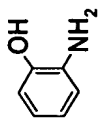
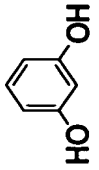
Structure	C12	C13	C14	C15	C16	C17	C18	C19	C20	C21	C22	C23	C24	C25	C26	C27	C28	C29
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
							X	X	X	X	X	X	X	X	X	X	X	X
																		
																		
																		
																		
							X									X		
	X	X	X	X	X	X		X								X	X	X

Table B (continued). Dye Combinations

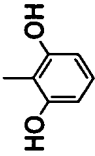
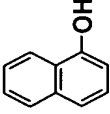
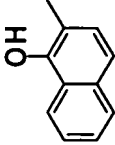
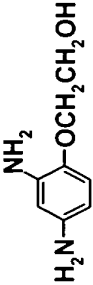
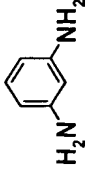
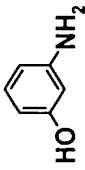
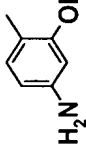
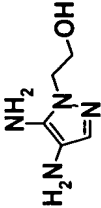
Structure	C12	C13	C14	C15	C16	C17	C18	C19	C20	C21	C22	C23	C24	C25	C26	C27	C28	C29
									X								X	
	X									X								X
		X									X							
			X									X						
				X									X					
					X									X				
						X									X			
																		

Table C. Dye Combinations

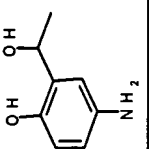
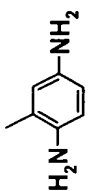
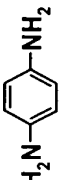

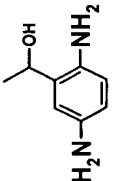
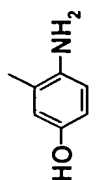
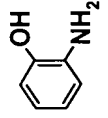
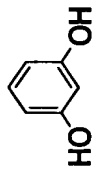
Structure	C30	C31	C32	C33	C34	C35	C36	C37	C38	C39	C40	C41	C42	C43	C44	C45	C46	C47
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
	X	X	X	X	X													
							X	X	X	X	X	X	X	X	X		X	X
																		
																		
																		
						X									X			
	X	X	X	X	X													X

Table C (continued). Dye Combinations

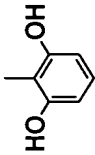
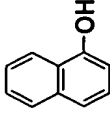
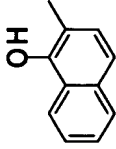
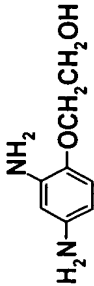
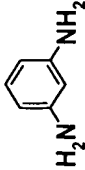
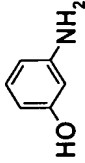
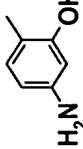
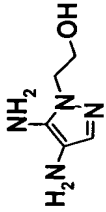
Structure	C30	C31	C32	C33	C34	C35	C36	C37	C38	C39	C40	C41	C42	C43	C44	C45	C46	C47
								X								X		
									X								X	
	X									X								X
		X									X							
			X									X						
				X									X					
					X									X				
																		

Table D. Dye Combinations

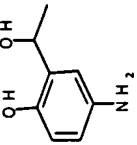
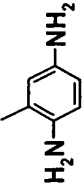


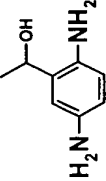
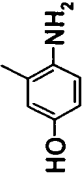
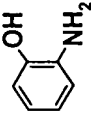
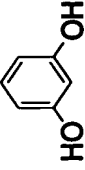
Structure	C48	C49	C50	C51	C52	C53	C54	C55	C56	C57	C58	C59	C60	C61	C62	C63	C64	C65
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
																		
	X	X	X															
																		
					X	X	X	X	X	X	X	X	X	X	X	X	X	X
																		
					X									X				
	X	X	X	X		X								X	X	X	X	X

Table D (continued). Dye Combinations

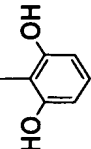
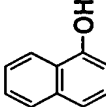
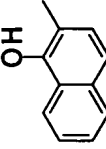
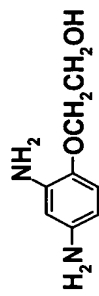
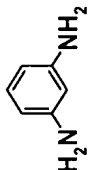
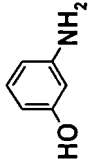
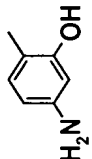
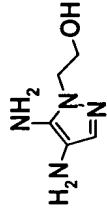
Structure	C48	C49	C50	C51	C52	C53	C54	C55	C56	C57	C58	C59	C60	C61	C62	C63	C64	C65
							X								X			
								X								X		
									X								X	
	X									X								X
		X									X							
			X									X						
				X									X					
																		

Table E. Dye Combinations

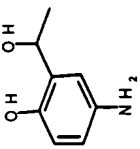
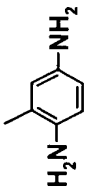
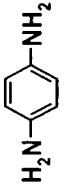

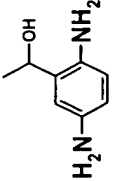
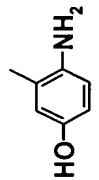
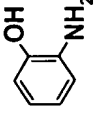
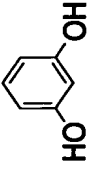
Structure	C66	C67	C68	C69	C70	C71	C72	C73	C74	C75	C76	C77	C78	C79	C80	C81	C82	C83
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
																		
																		
																		
	X	X	X															
																		
				X									X					
	X	X	X		X													X

Table E (continued). Dye Combinations

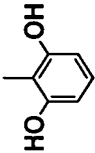
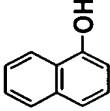
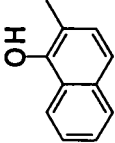
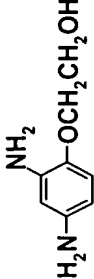
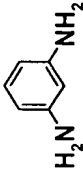
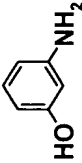
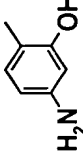
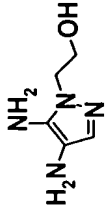
Structure	C66	C67	C68	C69	C70	C71	C72	C73	C74	C75	C76	C77	C78	C79	C80	C81	C82	C83
						X								X				
							X								X			
								X								X		
									X								X	
	X									X								X
		X									X							
			X															
				X														X

Table F. Dye Combinations

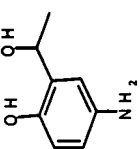
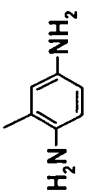
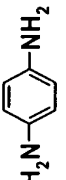

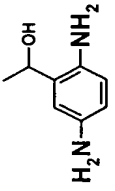
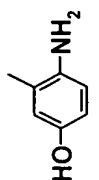
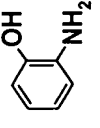
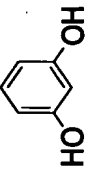
Structure	C84	C85	C86	C87	C88	C89	C90	C91	C92	C93	C94	C95	C96	C97	C98	C99	C100	C101
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
			X	X	X	X	X	X	X	X	X							
														X	X	X	X	X
																		
																		
																		
			X															
	X	X		X														

Table F (continued). Dye Combinations

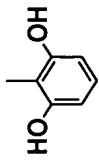
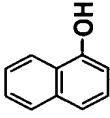
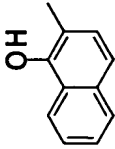
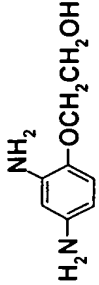
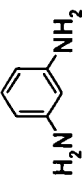
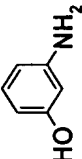
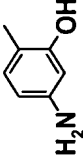
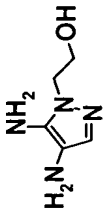
Structure	C84	C85	C86	C87	C88	C89	C90	C91	C92	C93	C94	C95	C96	C97	C98	C99	C100	C101
					X									X				
						X									X			
							X									X		
								X									X	
									X									X
	X									X								
		X									X							
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

Table G. Dye Combinations

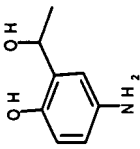
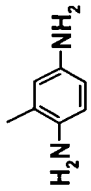
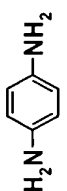

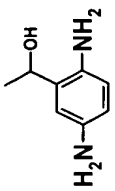
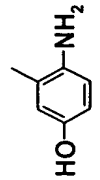
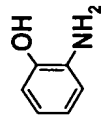
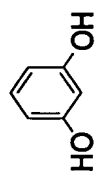
Structure	C102	C103	C104	C105	C106	C107	C108	C109	C110	C111	C112	C113	C114	C115	C116	C117	C118	C119
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
												X	X	X	X	X	X	X
	X	X																
																		
			X	X	X	X	X	X	X	X	X							
																		
			X									X						
				X								X	X	X	X	X	X	X

Table G (continued). Dye Combinations

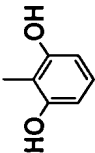
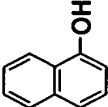
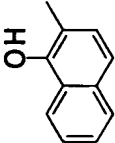
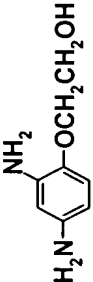
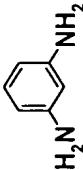
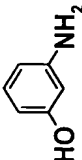
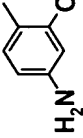
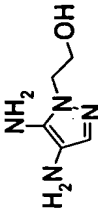
Structure	C102	C103	C104	C105	C106	C107	C108	C109	C110	C111	C112	C113	C114	C115	C116	C117	C118	C119
					X								X					
						X								X				
							X								X			
								X								X		
									X								X	
	X									X								X
		X									X							
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

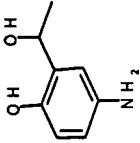
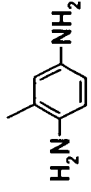
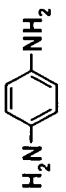

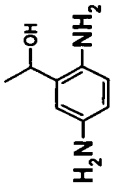
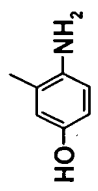
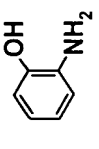
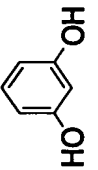
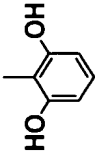
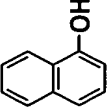
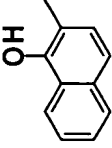
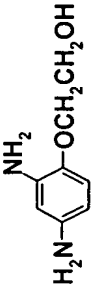
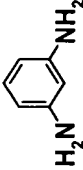
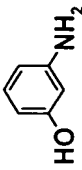
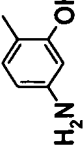
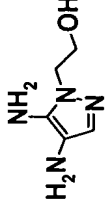
Table H. Dye Combinations																	
Structure	C120	C121	C122	C123	C124	C125	C126	C127	C128	C129	C130	C131	C132	C133	C134	C135	C136
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
	X																
		X	X	X	X	X	X	X	X								
																	
										X	X	X	X	X	X	X	X
																	
		X								X							
	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

Table H (continued). Dye Combinations

Structure	C120	C121	C122	C123	C124	C125	C126	C127	C128	C129	C130	C131	C132	C133	C134	C135	C136
			X								X						
				X								X					
					X								X				
						X								X			
							X								X		
								X								X	
									X								X
	X																
	X																

With the foregoing description of the invention, those skilled in the art will appreciate that modifications may be made to the invention without departing from the spirit thereof. Therefore, it is not intended that the scope of the invention be limited to the specific embodiments illustrated and described.

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